Dated: April 1, 1996.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 96–8529 Filed 4–5–96; 8:45 am]

BILLING CODE 4410–09–M

Importation of Controlled Substances; Notice of Application

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with § 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on February 22, 1996, Stepan Company, Natural Products Department, 100 W. Hunter Avenue, Maywood, New Jersey 07607, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

	Schedule
Drug:	
Coca Leaves (9040)	II
Cocaine (9041)	II
Benzoylecgonine (9180)	II

The firm plans to import the listed controlled substances to manufacture bulk controlled substances.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.54 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (30 days from publication).

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1311.42 (b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import basic classes of any controlled substances in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1311.42 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: April 1, 1996.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 96–8530 Filed 4–5–96; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substance; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on February 22, 1996, Stepan Company, Natural Products Department, 100 W. Hunter Avenue, Maywood, New Jersey 07607, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

	Schedule
Drug:	
Cocaine (9041)	II
Benzoylecgonine (9180)	II

The firm plans to manufacture the listed controlled substances in bulk for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the above application.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than June 7, 1996.

Dated: April 1, 1996.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 96–8531 Filed 4–5–96; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF LABOR

Occupational Safety and Health Administration

Maritime Advisory Committee for Occupational Safety and Health: Appointment of Members

AGENCY: Occupational Safety and Health Administration (OSHA), U.S. Department of Labor.

ACTION: Notice of appointment of members to the Maritime Advisory Committee for Occupational Safety and Health (MACOSH).

SUMMARY: The Secretary of Labor has established an advisory committee to advise the Assistant Secretary for the Occupational Safety and Health Administration (OSHA) on issues relating to the delivery of occupational safety and health programs, policies, and standards in the maritime industries of the United States. The committee will provide a collective expertise not otherwise available to the Secretary to address the complex and sensitive issues involved. Committee members have been appointed from government agencies, the shipbuilding industries and longshoring, labor and professional associations.

ADDRESSES: Any written comments in response to this notice should be sent to the following address: OSHA, Office of Maritime Standards, Room N–3621, 200 Constitution Avenue, NW., Washington, DC 20210. Phone (202) 219–7234, fax (202) 219–7477.

FOR FURTHER INFORMATION CONTACT: Mr. Larry Liberatore, Office of Maritime Standards, OSHA, (202) 219–7234.

SUPPLEMENTARY INFORMATION: MACOSH is intended to address the concerns of the entire maritime community, focusing on the shipyard and marine cargo (longshoring) handling industries. This committee will continue the efforts of the previously chartered Maritime Advisory Committee for Occupational Safety and Health. MACOSH is consistent with the President's initiative to make the U.S. shipyard industry competitive in the worldwide community. Furthermore, MACOSH will be able to focus on the resolution of controversial issues, particularly those with international implications,